## **AMENDMENTS TO THE CLAIMS**

Claims 1-34 (Canceled)

- 35. (Currently amended) A transgenic mouse comprising a disruption in an endogenous intestinal alkaline phosphatase gene, wherein where the disruption is homozygous, the transgenic mouse <u>lacks production of functional intestinal alkaline phosphatase protein and exhibits</u>, relative to a wild-type mouse, a nociceptive abnormality or abnormal activity level.
- 36. (Canceled)
- 37. (Canceled)
- 38. (Previously added) The transgenic mouse of claim 35, wherein the abnormal activity level comprises decreased activity.
- 39. (Previously added) The transgenic mouse of claim 38, wherein the decreased activity is characterized by a decreased velocity during ambulatory episodes in an open field test.
- 40. (Currently amended) A method of producing a transgenic mouse comprising a disruption in an endogenous intestinal alkaline phosphatase gene, the method comprising:
  - (a) introducing an intestinal alkaline phosphatase gene targeting vector into a murine embryonic stem cell;
  - (b) introducing the cell into a blastocyst;
  - (c) implanting the resulting blastocyst into a pseudopregnant mouse, wherein said pseudopregnant mouse gives birth to a chimeric mouse; and
  - (d) breeding the chimeric mouse to produce the transgenic <u>mouse</u>, wherein where the disruption is homozygous, the mouse <u>lacks production of functional</u> <u>intestinal alkaline phosphatase protein and exhibits a nociceptive abnormality or an</u> activity level abnormality, relative to a wild-type mouse.
- 41. (Previously added) A cell obtained from the transgenic mouse of claim 35.

Claims 42-46 (Canceled)